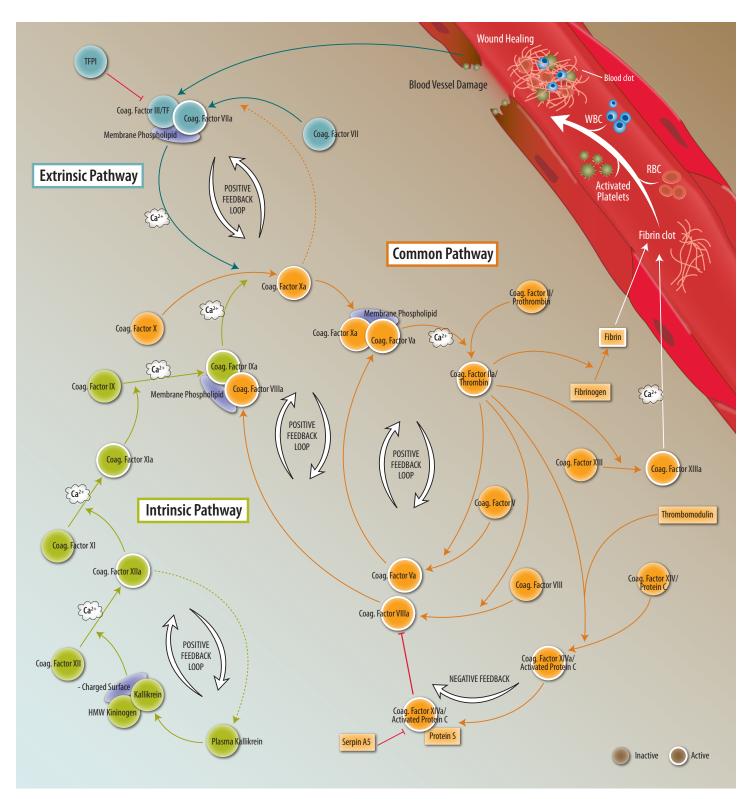


## **Blood Coagulation**



The Blood Coagulation Cascade Promotes the Formation of a Fibrin Clot. Following damage of a blood vessel, the extrinsic pathway of coagulation (blue arrows) is initiated by Coagulation Factor III/Tissue Factor (TF) which forms a complex with Coagulation Factor VIIa and phospholipid, in the presence of Ca<sup>2+</sup>, to activate Coagulation Factor Xa and rapidly generate Thrombin (IIa). Thrombin cleaves Fibrinogen to produce Fibrin which polymerizes in the presence of Coagulation Factor XIII to form a Fibrin clot. The slower, intrinsic pathway of coagulation (green arrows) provides an alternate mechanism for activation of Coagulation Factor Xa. It is initiated by Coagulation Factor XII, Plasma Kallikrein, and high molecular weight Kininogen binding to damaged subendothelial tissue. This results in the cleavage and activation of Coagulation Factor XIIa, which activates Coagulation Factor XIa. This factor the cleaves and activates Coagulation Factor IXa. Coagulation Factor Xa, and cagulation Factor VIIIa, activates Coagulation Factor Xa. Events downstream of the activation of Factor Xa are common to both the intrinsic pathways of coagulation (orange arrows). Activated Factor Xa cleaves Prothrombin to generate active Thrombin (IIa) which can then cleave Fibrinogen to produce Fibrin onomers. These monomers are cross-linked by Factor XIII to form a Fibrin clot.

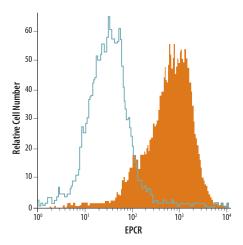
## **Blood Coagulation**

Injuries that damage blood vessels promote blood coagulation, a rapid response to initiate hemostasis and protect the host from excessive blood loss. Blood coagulation results from a series of proteolytic reactions involving the step-wise activation of coagulation factors. Subsets of these factors can be activated by two distinct pathways, the extrinsic, or tissue damage pathway (blue arrows), and the intrinsic, or contact pathway (green arrows). Each is initiated by different factors, but both converge upon a single common pathway (orange arrows) that leads to the activation of Coagulation Factor Xa, and the conversion of Prothrombin/Coagulation Factor II to active Thrombin/Coagulation Factor IIa. Thrombin converts Fibrinogen to Fibrin monomers which polymerize to form a Fibrin clot. The Fibrin clot acts in concert with activated platelets at the site of the injury to form a blood clot that stabilizes the damaged tissue and prevents further blood loss.

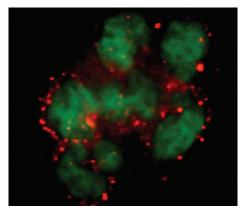
In addition to directly generating active Fibrin, Thrombin activates Coagulation Factor XIII, which stabilizes Fibrin and promotes its polymerization. Thrombin also activates Coagulation Factors V, VIII, and Protein C. These factors enhance or inhibit Thrombin production through positive or negative feedback. Factors Va and VIIIa promote Thrombin production by positively regulating either the cleavage of Prothrombin itself, or the cleavage and activation of Coagulation Factor Xa, respectively. In contrast, activation of Protein C by Thrombin binding to Thrombomodulin leads to the degradation of Factors Va and VIIIa, and inhibits the cleavage of Prothrombin. These forms of feedback regulation, along with the sequential activation of clotting factors, allow precise control of the blood coagulation cascade. This tight regulation is critical to prevent excessive blood loss associated with too little clotting, or too much clotting, which could result in the blockage of a blood vessel and lead to a heart attack or a stroke. Identifying other regulatory mechanisms may reveal additional molecular targets for exogenous control of clotting activity. R&D Systems offers a variety of research reagents useful for the characterization of molecules involved in blood coagulation pathways.

Molecule	Recombinant & Natural Proteins	Antibodies	ELISAs/ Assays	Molecule	Recombinant & Natural Proteins	Antibodies	ELISAs/ Assays
ADAMTS13	Н	н	н	Kininogen	НМ	НМ	
Angiostatin	Н	Н		α2-Macroglobulin	Н	НМ	н
Annexin A2		HMR	Н	PAR1	н	н	
Annexin A6	Н	н		PAR2		н	
Annexin V	Н	н	Ms	Plasma Kallikrein/KLKB1	НМ	НМ	
Apolipoprotein H/ApoH	НМ	HMR		Plasminogen	н	н	
Coagulation Factor II/Thrombin	Н	Н		Plasminogen Kringle 5		М	
Coagulation Factor III/Tissue Factor	НМ	НМ	Н	Protein S/PROS1		НМ	
Coagulation Factor VII	НМ	НМ		SDNSF/MCFD2		НМ	
Coagulation Factor IX coming soon!	н			Serpin A1/a1-Antitrypsin	НМ	НМ	н
Coagulation Factor X	НМ	н		Serpin A5	н	НМ	
Coagulation Factor XI	НМ	Н		Serpin C1/Antithrombin-III	Н	НМ	н
Coagulation Factor XIV/Protein C	НМ	НМ		Serpin D1/Heparin Cofactor II	НМ	НМ	
Collagen I	BR		Ms	Serpin E1/PAI-1	н	НМ	Н
Collagen II	Н	M	Н	TFPI	НМ	НМ	НМ
Collagen IV	M	н	Ms	TFPI-2	н	НМ	
Collagen XXV $\alpha$ 1		н		Thrombomodulin/BDCA-3	НМ	НМ	НМ
EPCR	М	НМ	Н	u-Plasminogen Activator/Urokinase	Н	Н	Н
Glycoprotein V/CD42d	НМ	НМ		uPAR	НМ	НМ	НМ
CD42b/GP1ba	НМ	н		vWF-A2	Н	Н	Н
GPVI	НМ	НМ		KEY: H: Human M: Mouse R: Rat B: Bovine Ms	: Multi-species		

## Blood Coagulation Research Reagents Available from R&D Systems



Detection of Endothelial Protein C Receptor (EPCR) in Mouse bEnd-3 Cells by Flow Cytometry. Mouse bEnd-3 endothelial cells were stained with biotinylated anti-mouse EPCR polyclonal antibody (Catalog # BAF2749; filled histogram) or biotinylated normal goat IgG (Catalog # BAF108; open histogram) followed by APC-conjugated streptavidin (Catalog # F0050).



**Detection of Coagulation Factor VII in Human Peripheral** Blood Mononuclear Cells (PBMC). Coagulation Factor VII was detected in PHA-stimulated human PBMC using antihuman Coagulation Factor VII monoclonal antibody (Catalog # MAB2338). Cells were stained using a Rhodamine Red™ X-conjugated anti-mouse IgG secondary antibody (red) and counterstained with FluoroNissI™ Green (green).

## **R** Systems







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